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AMENDMENTS TO THE CLAIMS

- Claim 1. (Currently Amended) A method of qualifying the prostate cancer status in a subject comprising:
- (a) measuring the amount of at least one-biomarker in a sample from the subject, wherein the biomarker is selected from the group consisting of

	Marker I:	having a molecular weight of about 7.808 kD in a biological	
sample	from the subje	ect.	
	Marker II:	having a molecular-weight of about 14,576 kD	

Marker II:	having a molecular-weight of about 14.576 kD
Marker III:	having a molecular weight of about 2.062 kD
	having a molecular weight of about 7.974 kD
	having a molecular weight of about 6.677-kD
	having a molecular weight of about 3.936 kD
	having a molecular weight of about 60.958 kD
Marker-VIII:-	having a molecular weight of about 5.149 kD
	having a molecular weight of about 5.861-kD
	having a molecular weight of about 28.098 kD
Marker XI:	- having a molecular weight of about 2.996 kD
	having a molecular weight of about 24.346 kD
	having a molecular weight of about 6.722 kD
	having a molecular weight of about 5.999 kD
- Marker XV:	having a molecular weight of about 6.158 kD
	having a molecular weight of about 55.785 kD
Marker XVII:	having a molecular weight of about 2.540 kD
	having a molecular weight of about 8.019 kD
	having a molecular weight of about 4.658 kD
Marker XX:	having a molecular-weight of about 14.703 kD
Marker XXI:	-having a molecular-weight of about 2.68 kD
Marker XXX:	having a molecular weight of about 3.16 kD
Marker XXIII	having a molecular weight of about 10.3 kD
Marker XXIV	thaving a molecular weight of about 10.8 kD

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Marker XXV: having a molecular weight of about 12.7 kD

Marker XXVI:having a molecular weight of about 17.9 kD

Marker XXVII: having a molecular weight of about 2.79 kD

Marker XXVIII: having a molecular weight of about 3.32 kD

Marker XXIX: having a molecular weight of about 4.29 kD

Marker XXX: having a molecular weight of about 15.9 kD

Marker XXXI: having a molecular weight of about 16.1 kD

Marker XXXII: having a molecular weight of about 16.3 kD, and combinations thereof, and

(b) wherein a decrease in the amount of the marker as compared to a control is indicative that the subject has correlating the measurement with prostate cancer-status.

- Claim 2, (original) The method of claim I further comprising:
- (c) managing subject treatment based on the status.
- Claim 3. (original) The method of claim 2, wherein managing subject treatment is selected from ordering more tests, performing surgery, and taking no further action.
 - Claim 4. (Currently Amended) The method of claim 2 further comprising:

 (d) measuring the at least one biomarker after subject management.

Claims 5-10. (Cancelled)

- Claim 11. (currently amended) The method of <u>claim 1</u> any of claims 1-10 wherein the marker is detected by mass spectrometry.
- Claim 12. (currently amended) The method of <u>claim 1</u> any of claims 1 10 wherein the marker is detected by capturing the marker on a biochip having an affinity surface and detecting the captured marker by SELDI.

Claims 13-77. (cancelled)

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